

I023390

Bayer CropScience



October 27, 2011

Document Processing Desk 6(a)(2)
Office of Pesticide Programs (7504P)
U. S. Environmental Protection Agency
Room S-4900, One Potomac Yard
2777 South Crystal Drive
Arlington, VA 22202-4501

RE: 6(a)(2) Incidents Accumulated for the Month of September 2011

Dear Sir/Madam:

Reportable incidents accumulated for the month of September 2011 for Bayer CropScience and Bayer Environmental Science are attached.

The information with this letter is being submitted to the EPA pursuant to the Agency's interpretation of requirements imposed on registrants by Section 6(a)(2) of FIFRA. This information does not necessarily constitute additional factual information regarding unreasonable adverse effects within the meaning of 6(a)(2). It is being submitted to enable the Agency to make its own assessment of the information.

If you have questions or concerns, please do not hesitate to contact me at any time.

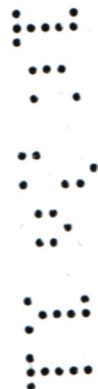
Sincerely,

Gerret Van Duyn
Compliance Manager
State Regulatory and Documentation Services
919-549-2914

CC: AE Coordinator, CA Department of Pesticide Regulation
Jeanine Broughel, NY Department of Environmental Conservation

/attachment

Bayer CropScience
RTP
P. O. Box 12014
RTP, NC 27709
Tel. 919 549-2000



Personal privacy information

-003

Row 1 Administrative Data	Reporter Name [REDACTED]	Submission date. 10/27/2011	Contact person (if different than reporter)	Internal ID 860707
	Address [REDACTED]		Address	
			Phone #	
	Incident Status: New	Location and date of incident Cynthiana, KY USA 08/04/2011	Date registrant became aware of incident. 09/29/2011	Was incident part of larger study? No
Row 2 Pesticide(s) Involved	EPA Registration # (Product 1) 72155-80		EPA Registration # (Product 2)	EPA Registration # (Product 3)
	A.I. (s) Beta-Cyfluthrin, sodium o-phenylphenate		A.I. (s)	A.I. (s)
	Product 1 name Home Pest plus Germ Killer Indoor & Outdoor Killer RTU (1 Gal)		Product 2 Name	Product 3 Name
	Exposed to concentrate prior to dilution? NA		Exposed to concentrate prior to dilution?	Exposed to concentrate prior to dilution?
	Formulation		Formulation	Formulation
Row 3 Incident Circumstances	Evidence label directions were not followed? Yes Intentional misuse? Yes	Incident site: (examples include home, yard, school, industrial, nursery/greenhouse, surface water, commercial turf, building/office, forest/woods, agricultural (specify crop) right-of-way (rail, utility, highway)). Own Residence	Situation (act of using product): (examples include mixing/loading, reentry, application, transportation, repair/ maintenance of application equipment, manufacturing/ formulating). See Incident Description Notes	
	Applicator certified? UNK			
	How exposed: (examples include direct contact with treated surface, ingestion, spill, drift, runoff) See Incident Description Notes			

Brief description of incident circumstances.

Billings, Sharon Sep 29 2011 11:27AM
Tfr from Kirstin.

Hx: Caller used product on mattress about 6 to 8 weeks ago and about 1 week later caller developed a rash with bumps all over his body. Caller saw HCP who prescribed topical steroid and sxs persist. Caller stated he was aware this was not a labeled use of the product. Caller is inquiring how to remove product from mattress.

A: This is a misuse of the product; dermal irritation can result from prolonged dermal exposure. We refer you to your HCP for continued care. We refer you to Customer Service for questions regarding removal of product from treated surface and we refer you to your HCP for continued care. If any new or unexpected symptoms develop, please contact us 24/7 and refer to your reference number so that we can advise on further treatment. Gave case#, cb prn.

LeMaster, Steve Sep 30 2011 9:38AM
notified

Demographic information: Age: 24 Year(s) Sex: Male Occupation (if relevant) NA	Exposure route: Dermal	Was adverse effect result of suicide/homicide or attempted suicide/homicide? No	Was protective clothing worn (specify)? None Reported
If female, pregnant? NA	Was exposure occupational? Not indicated If yes, days lost due to illness: NA	Time between exposure and onset of symptoms: 1 week or less	
Type of medical care sought: (examples include none, clinic, hospital emergency department, private physician, PCC, hospital inpatient). Private MD/DVM-treated & released	List signs/symptoms/adverse effects Dermatological-Hives/Welts Dermatological-Rash		If lab tests were performed, list test names and results (If available, submit reports) None Reported
Exposure data: NA Amount of pesticide: NA Exposure duration: Acute < 8hrs Patient weight: Unknown			
Human severity category: HC			
<p>This box can be used to provide any explanatory or qualifying information surrounding the incident. (add additional pages if necessary)</p> <div style="border: 1px solid black; height: 150px; width: 100%;"></div> <div style="border: 1px solid black; width: 150px; float: right; padding: 5px;"> Internal ID # 860707 </div> <div style="clear: both;"></div>			